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APPLICATION NO. FILING DATE 09/644,605 08/24/2000	FIRST NAMED INVENTOR Patrick Briot	ATTORNEY DOCKET NO.  PET-1875	CONFIRMATION NO.
23599 7590 02/07/2002 MILLEN, WHITE, ZELANO & I 2200 CLARENDON BLVD. SUITE 1400 ARLINGTON, VA 22201	BRANIGAN, P.C.	PREISCH, I	NADINE G  PAPER NUMBER

Please find below and/or attached an Office communication concerning this application or proceeding.

<u>.</u>		Application No.	Applicant(s)
		09/644,605	BRIOT ET AL.
	Office Action Summary	Examiner	Art Unit
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	Responsive to communication(s) filed	on <u>23 November 2001</u> .	
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2a)⊠ 3)□	This action is <b>FINAL</b> . 20 Since this application is in condition for closed in accordance with the practice	and formal m	atters, prosecution as to the ments is c.D. 11, 453 O.G. 213.
Disposit	ion of Claims		
4)⊠	ousim(a) 1.2 and 4-20 is/are pending i	n the application.	
	4a) Of the above claim(s) is/are	withdrawn from consideration.	
5)□	Claim(s) is/are allowed.		
6)⊠	Claim(s) 1,2 and 4-20 is/are rejected.		
<b>-</b> √□	claim(s) is/are objected to.		
8)	Claim(s) are subject to restrict	on and/or election requirement.	
Applica	tion Papers		
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10)	• - 1	-ITI cocontad of hill I objected to b	ny trie Examinion.
15)	The drawing(s) filed on is/are:  Applicant may not request that any obje	ection to the drawing(s) be held in ab	I disapproved by the Examiner.
11)	The proposed drawing correction filed	on is: a)[_] approved b)[	Tidisabbiosed by the Emmission
1	If approved, corrected drawings are rec	fulled in reply to this sines	
12)	The oath or declaration is objected to	by the Examiner.	
l .	·- a ac 440 and 120		- 0.440(a) (d) or (f)
13\0	y under 35 U.S.C. §§ 119 and 120  ☑ Acknowledgment is made of a claim	for foreign priority under 35 U.S	.C. 8 119(8)-(a) or (i).
10/2	NO All b) Some * c) None of:		
		documents have been received.	· Audication No
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	<ol> <li>Copies of the certified copies application from the Internal</li> </ol>	of the priority documents have be national Bureau (PCT Rule 17.2)	(a)).
	* See the attached detailed Office action	for domestic priority under 35 U.	S.C. § 119(e) (to a provisional application)
14)[	<ul><li>Acknowledgment is made of a claim</li><li>a) ☐ The translation of the foreign la</li></ul>	nguage provisional application h	as been received.
15)	a) $\square$ The translation of the foreign la $\square$ Acknowledgment is made of a claim	for domestic priority under 35 U	.S.C. §§ 120 and/or 121.
	ment(s)	4) 🔲 Inte	erview Summary (PTO-413) Paper No(s)
	Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review Information Disclosure Statement(s) (PTO-1449)	(PTO-948) 5) Not	ice of Informal Patent Application (PTO-152)
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### DETAILED ACTION

## Withdrawal of Objection to the Specification

Applicants' amendment filed 11-23-01 in paper no.8 is sufficient to overcome the previous objection to the abstract.

### Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

Claims 14, 16, and 17 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicants regard as the invention.

In claims 14, 16 and 17, it is unclear if the terminology in parenthesis is intended to be part of the claimed subject matter. If applicants intend to claim such subject matter, it is suggested that applicants' remove the parenthesis and insert the subject matter into the body of the claim.

In claims 14 and 16, applicants refer to both "VIB" and "VI" metals. It appears as "VI" should be changed to "VIB" to be consistent.

In claim 16, the terminology of "the phosphorous oxides" lacks antecedent basis in the claims because none of the preceding claims contain phosphorous oxide.

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### Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1-2, 4-5, and 7-18 are rejected under 35 U.S.C. 103(a) as being unpatentable over Billon (5,525,209) in view of Walker (3,507,786) and Cody et al.(5,911,874).

Applicants are claiming a process for producing a high viscosity index oil from a feed containing constituents boiling above 300°C. The process involves a) reacting a feed with a catalyst comprising at least one amorphous non-zeolitic matrix and at least one Group VIII or VIB metal or metal compound; b) fractionating the effluent from step a) to separate at least one oil residue comprising mainly constituents with viscosity indices higher than the feed; c) fractionating the residue by thermal diffusion into oil fraction with high viscosity indices. Applicants claim an additional embodiment wherein a portion of the effluent from step a) is brought into contact with a catalyst comprising at least one zeolite, at least one matrix and at least one Group VIII or VIB metal or metal compound prior to being sent to step b).

The reference of Billon et al.(5,525,209) teaches a process for producing an oil with a high viscosity index between 95 and 150. See column 1, lines 8-11 and column 4, lines 5-10.

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The process comprises contacting a feed boiling above the temperature above 380°C with a catalyst in the presence of hydrogen. See column 1, lines 42-50. The catalyst contains an amorphous support and at least one Group VIII or VI element. See column 1, lines 48-50. Suitable catalyst supports include alumina, silica, silica-alumina, magnesia and clay. See column 2, lines 5-9. The catalyst further comprises a concentration of VIII or VI metals of 5-40% by weight expressed as the oxides with the weight ratio of the metal expressed as the oxides between Group VIB and VIII between 20 and 1.25. See column 2, lines 22-30. The concentration of phosphorous oxide is less than 15% by weight. See column 3, lines 1-5. First step process conditions include a temperature between 350°C and 430°, a pressure of 2 to 20 Mpa, a space velocity of 0.1 to 5hr<sup>-1</sup> and a hydrogen/hydrocarbon ratio of 150 to 2000 by volume. See column 2, lines 39-44 and column 1, lines 52-55.

The reference further teaches that the product obtained from the first step is passed to a second step wherein it is contacted with a second catalyst in the presence of hydrogen. See column 2, lines 49-54. The second catalyst contains a zeolite, a Group VI or VIII metal and a matrix (support). See column 2, lines 57-61. Billon et al.(5,525,209) teaches that the second step catalyst can comprise an acid HY having a SiO<sub>2</sub>/Al<sub>2</sub>O<sub>3</sub> molar ratio of 12-40. The reference further teaches that the zeolite has a unit cell size of 24.55 x 10<sup>-10</sup>m to 24.24 x 10<sup>-10</sup>m, a C<sub>Na</sub> greater than 0.85, a specific surface area of greater than 400 m<sup>2</sup>/g, a water vapor adsorption capacity at 25°C of 2.5 torr of greater than 6%, and a pore distribution comprising between 1% and 20% of the pore volume contained in pores with a diameter between 20 x 10<sup>-10</sup> and 80 x 10<sup>-10</sup> m. See column 3, lines 25-40. The reference also discloses that the zeolite is calcined at 1100°C. See column 3, lines 23-25. The weight of the zeolite with respect to the catalyst is 2 to

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80 wt%. See column 3, lines 10-14. Second step process conditions include a temperature between 350°C and 430°, a pressure of 2 to 20 Mpa, and a space velocity of 0.1 to 5hr<sup>-1</sup>. See column 3, lines 43-48.

Billon et al.(5,525,209) teaches that the product from the second step is fractionated to obtain a residue containing the oil and middle distillates. See column 3, lines 10-24. A portion of the residue can be recycled. See column 4, lines 14-17.

The reference of Billon et al.(5,525,209) succeeds in disclosing a process for the production of a high viscosity oil. In addition, the reference succeeds at disclosing a process with steps corresponding to applicants' initial hydrogen/catalyst contact, optional secondary catalyst/hydrogen contact, fractionation to obtain a residue and distillates and product recycle. Billon et al.(5,525,209) also discloses applicants' claimed process conditions and catalyst compositions.

Several differences are noted between the reference of Billon et al.(5,525,209) and applicants' claimed invention. The reference does not disclose the use of thermal diffusion to fractionate the residue. Also, the reference is silent about including a vapor/liquid step after the first processing step.

The reference of Walker (3,507,786) is cited to illustrate that it is known in the art that thermal diffusion can be used to separate hydrocarbon oils by their molecular shapes. See column 1, lines 56-61. The reference illustrates that thermal diffusion techniques are known to obtain different pour point/ V.I. fractions. See column 2, lines 60-72 and 1-15.

The reference of Cody et al.(5,911,874) is cited to show that it is conventional to separate undesirable lower boiling point products such as hydrogen sulfide and ammonia (e.g. gas) from

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the first stage conversion of product in a two stage hydroconversion process. See column 6, lines 13-21.

It would have been obvious to one of ordinary skill in the art at the time the invention was made desiring to separate the final residue obtained in the Billon et al.(5,525,209) process into selected VI and/or pour point fractions to include an additional thermal diffusion fractionation step because the reference of Walker (3,507,786) illustrates that it is within the level of ordinary skill in the art to separate desired V.I. and/or pour point fractions by thermal diffusion. In addition, it is within the level of ordinary skill to operate the thermal diffusion process at any temperature required to obtain desired product cuts. Applicants have not shown anything unexpected by utilizing a conventional thermal diffusion fractionation step to obtain desired product cuts.

In addition, applicants' reactor dimensions in claim 13 are not considered to be patentable distinctions because they are apparatus limitations which do not limit the process in a manipulative sense. It would have been obvious to one of ordinary skill in the art at the time the invention was made to select any thermal diffusion column dimensions that would accomplish a desired separation in the process encompassed by the modified teachings of Billon et al.(5,525,209), including the specific dimensions claimed by applicants, because it has been held that invention in a method must be found in the steps performed and not the apparatus employed. Ex Parte Hart 117 USPQ 193 (Bd PatApp & Int 1958).

It would have been obvious to one of ordinary skill in the art at the time the invention was made desiring to remove undesirable gaseous products such as hydrogen sulfide and ammonia produced in the first stage conversion of Billon et al.(5,525,209) to include a vapor

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separation step to remove such components because the reference of Cody et al.(5,911,874) illustrates that it is conventional to separate gaseous components such as hydrogen sulfide and ammonia in two stage hydroprocessing. Applicants have not shown anything unexpected by including a known separation step in the process of Billon.

### Claim Rejections - 35 USC § 103

Claims 6 and 19-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Billon (5,525,209) in view of Walker (3,507,786) as applied to claims 1-2, 4-5, and 7-18 above, and further in view of Garwood et al.(4,975,177).

A difference is noted between the process of Billon et al.(5,525,209) and applicants' claimed invention. The reference of Billon et al.(5,525,209) does not disclose dewaxing the final residue.

The reference of Garwood et al.(4,975,177) is cited to illustrate the general concept that an additional selective dewaxing step can be used to obtain a "target" pour point. See abstract, lines 4-6. The purpose of selective dewaxing is to remove undesirable waxy components which contribute to an unfavorable pour point without removing the desirable iso-paraffin components which contribute to high VI. See column 10, lines 50-60. Shape selective catalysts including ferrierite can be employed in the selective dewaxing. See column 11, lines 10-11.

Garwood et al.(4,975,177)'s disclosure of ferrierite is considered to encompass applicants' bridge distance limitations because applicants' specifically disclose that ferrierites meet the criteria of the molecular sieve (See applicants' spec, page 8, lines 20-23) used in the claimed invention.

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It would have been obvious to one of ordinary skill in the art at the time the invention was made desiring to improve the pour point of the high viscosity fraction to include an additional dewaxing step because the reference of Garwood et al.(4,975,177) teaches that a selective dewaxing functions to desirably improve the pour point while maintaining VI.

Applicants have not shown anything unexpected by including an additional known prior art step.

In addition, it would have been obvious to one of ordinary skill in the art at the time the invention was made to select a dewaxing composition with applicants' bridge distance because Garwood et al. (4,975,177) illustrates that it is conventional to employ a ferrierite which is considered to encompass applicants' claimed bridge distance.

#### Response to Arguments

Applicants' arguments filed 11-23-01 in paper no.8 have been fully considered but they are not persuasive.

Applicants' arguments against the combination of Billion and Walker or Cody are not persuasive in overcoming the rejections. Applicants' arguments are considered to be a piecemeal attack of the references. It is maintained that the rejection is proper because the "combination" of references suggests the limitations in applicants' claims. For instance, the secondary reference of Cody is relied on to illustrate the conventionality of removing undesirable low boiling point gases such as hydrogen sulfide and ammonia from a hydrocarbon product. The fact that Cody discloses other steps that are not found in the primary reference does not diminish its illustration of the conventionality of removing undesirable gas components from a hydrotreated product. One of ordinary skill in the art would be motivated to modify the process of Billion to

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remove such components from an effluent obtained from any hydrotreating step since they are undesirable.

Furthermore, newly submitted claim 18 is considered to been encompassed by the applied art because the claim does not "exclude" the second treatment disclosed by the primary reference.

Applicants' amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Nadine Preisch whose telephone number is 703-305-2667. The examiner can normally be reached on Monday through Thursday from 7:30 am to 6:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Marian Knode can be reached on 703-308-4311. The fax phone numbers for the organization where this application or proceeding is assigned are 703-305-3599 for regular communications and 703-305-5408 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 308-0661.

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N.P.

February 6, 2002

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